EXHIBIT 67

U. S. Departr t of Justice
Drug Enforcement Administration
Office of Training
P.O. Box 1475
Quantico, Virginia 22134-1475

JAN 12

Mr. Steve Mays
Manager, Regulatory Affairs
Amerisource Bergen
P.O. Box 959
Valley Forge, Pennsylvania 19482

Dear Mr. Mays:

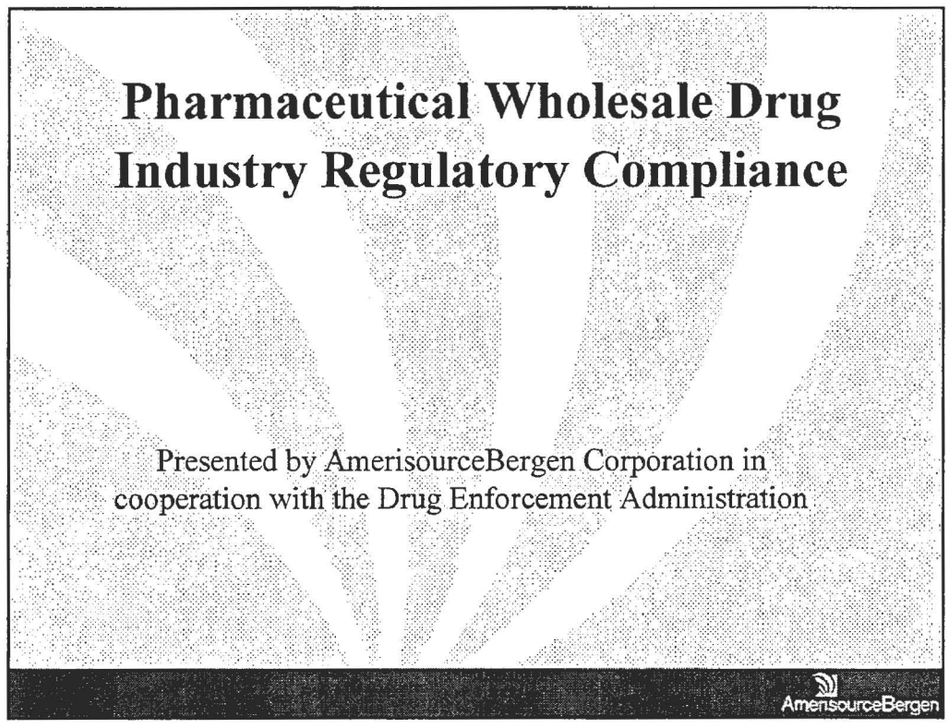
This letter is to confirm previous arrangements made by the Drug Enforcement Administration (DEA) Office of Training Acting Unit Chief Tom Prevoznik for a tour of the Bergen Brunswig facility in Richmond, Virginia, by our Diversion Investigator Trainees. I appreciate your cooperation, and I am certain that the visit to your distribution plant will be a valuable learning experience for our students.

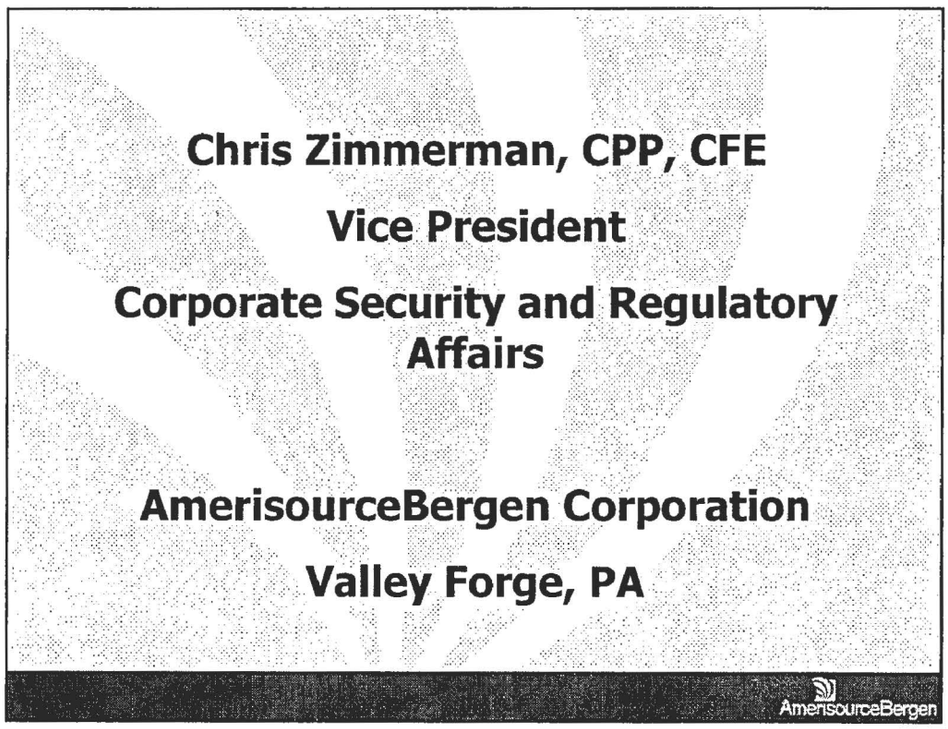
The visit to Bergen Brunswig is scheduled for Wednesday, March 9, 2005. Approximately 40 employees participating in the tour will be arriving by bus at approximately 9 a. m., and we will depart your facility at approximately 12 noon for the return trip to Quantico, Virginia.

Again, thank you for your cooperation and support of the DEA Office of Training. Mr. Prevoznik will be in contact with you regarding further arrangements. In the meantime, if you should have any questions, please do not hesitate to contact him at (703) 632-5217.

Sincerely,

John R. McCarty
Special Agent in Charge





Statement of Goals

- The goal of this program is to provide the participants with an overview of the Pharmaceutical (Drug) Wholesale industry and the Wholesaler's compliance with 21 C.F.R. 1300 to the End. In addition, we will provide examples and methods of standard operating procedures of a full-line pharmaceutical wholesaler in an attempt to educate and thus enhance and build on the good working relationship between the Industry and DEA.

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Objectives

At the end of the presentation attendees should:

- Understand the basic functions of a drug wholesaler with regard to distribution of controlled substances
- Be familiar with the basic structure and departments of a drug wholesaler
- Understand the Industry, basic operations and procedures and how they relate to 21 CFR 1300 to the end
- Evaluate and understand the different types of reports and documents available during an audit

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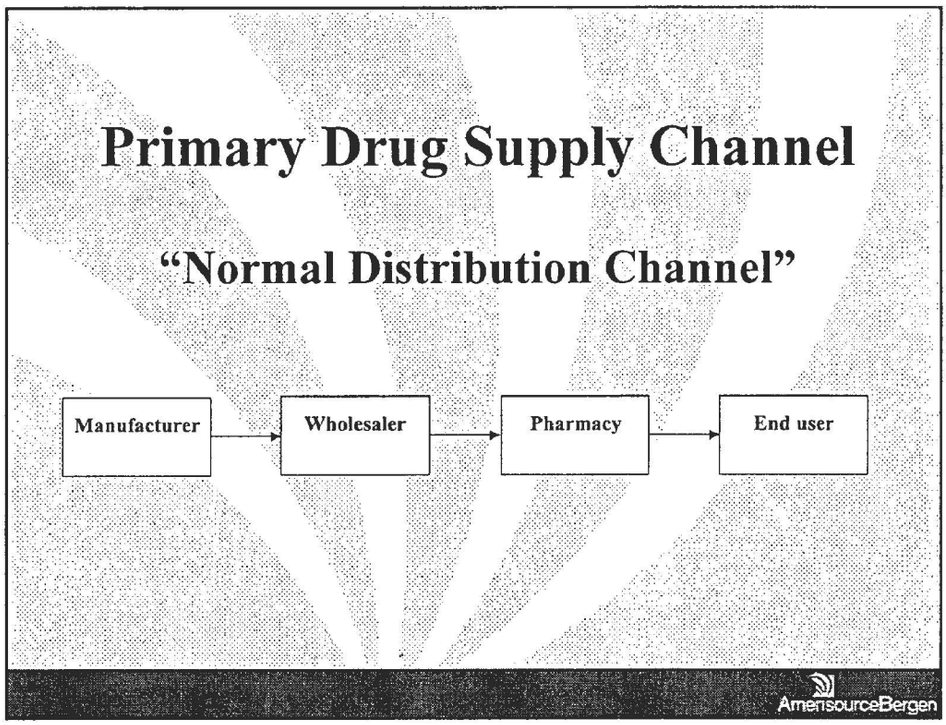

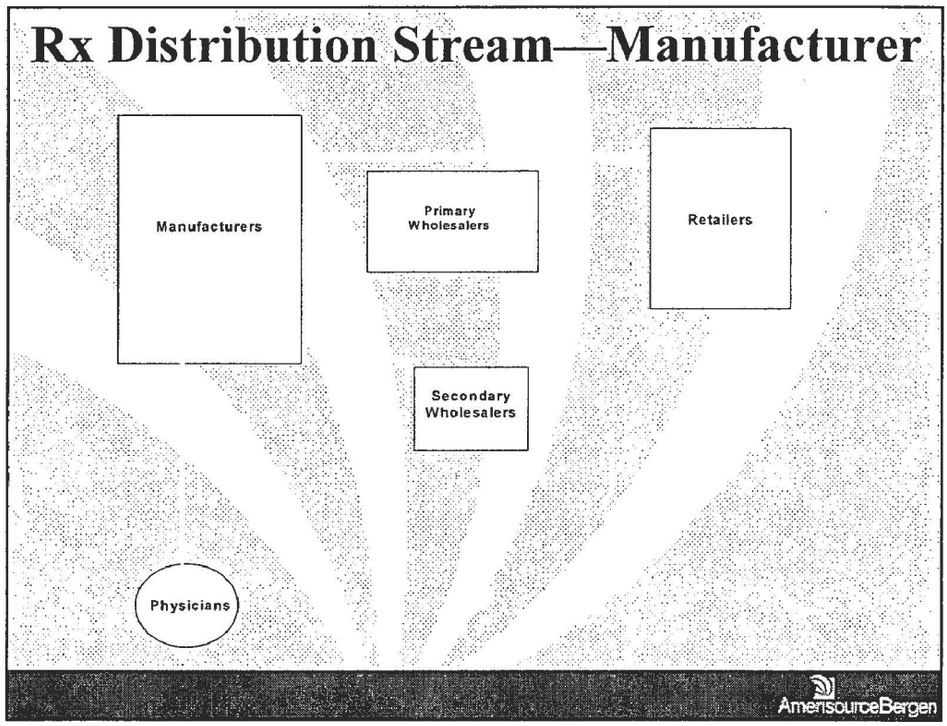

Wholesaler Definitions

- Primary Wholesaler
  - Purchase majority of their products from the manufacturer
  - Distribute the majority of their products to providers (Pharmacies, Hospitals, Clinics, etc.)
  - Approximately 90% of all drugs distributed are through the "Big 3" (ABC, Cardinal Health, McKesson)
- Secondary Wholesalers (NMV's, ASV's)
  - Purchase the majority of their products from other wholesalers
  - Distribute the majority of their product primarily to other wholesalers
  - There are approximately 7,000 licensed wholesalers in the United States (50 primary wholesalers)

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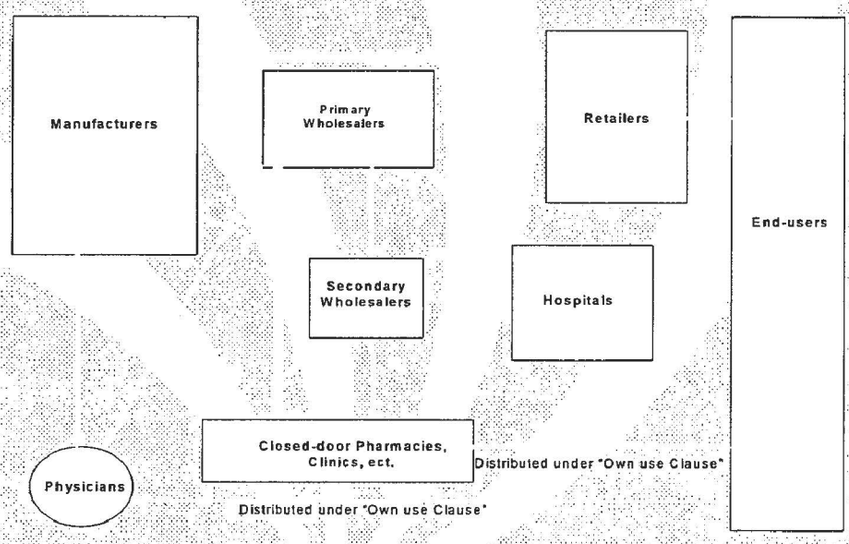
Rx Distribution Stream



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Wholesale Drug Distributors

- There are over 3,000 different manufacturers of pharmaceuticals and health care products in the United States
  - ABC maintains approximately 140,000 SKUs (stock keeping units)
- There are over 130,000 pharmacy sites in the U.S.
  - Retail
  - Closed Door
  - Hospital

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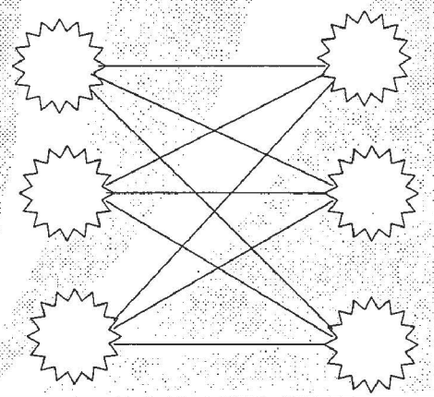
Role of the Wholesale Distributor

Manufacturers Pharmacies

Without Distributors

Confusion and Complexity

If all 130,000 pharmacy sites ordered product from all 3,000 manufacturers once a month the industry would generate 4.9 billion transactions each year



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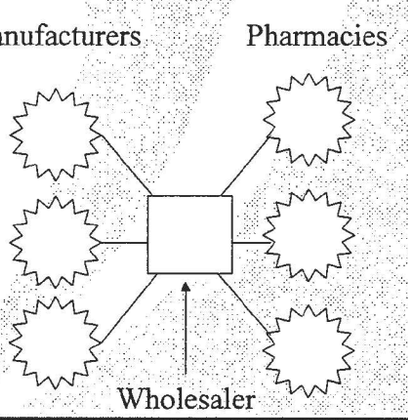
Role of the Wholesale Distributor

With Distributors

Control and Common Sense

Utilizing the 50 primary drug wholesalers that number is cut to 80 million transactions (99% reduction)



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Wholesale Drug Distributors

- There are approximately 50 primary wholesalers who operate approximately 230 distribution centers nationwide which generate approximately $120 billion in sales annually
- The top 3 wholesalers account for approximately 90% of the market
- In the past decade the industry has cut operating expenses by over 30 percent
- Cutting operating expenses has not curved the steady decline in profit margins
- Average product mark-up is 1%
- Estimated net profits for Wholesale Drug Distributors is less than 1 percent (0.65%) of sales

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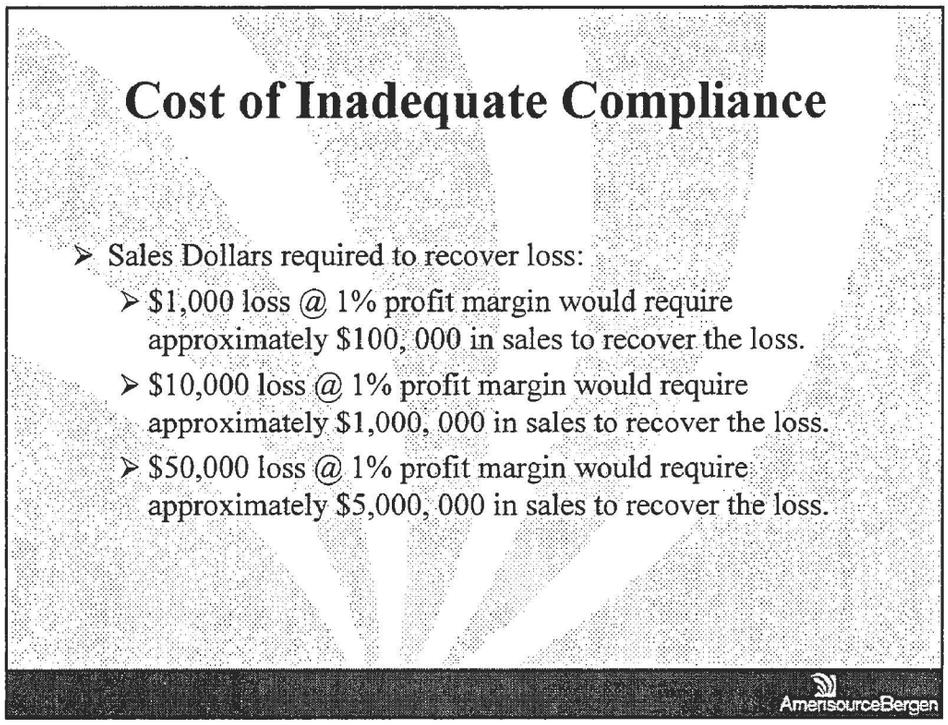

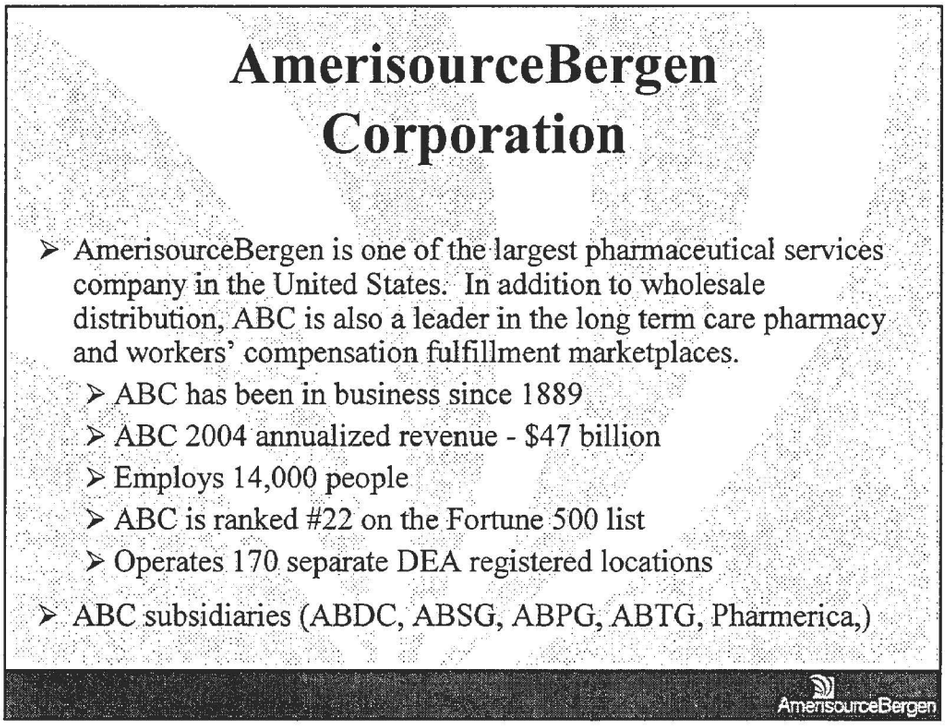

AmerisourceBergen Commitment to Compliance

- Regulatory Compliance is driven operationally on a daily basis at each individual distribution center with oversight, guidance, and assurance from the corporate office.
- Six-Hour Security & Regulatory Compliance Training Program for "compliance critical" Associates.

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AmerisourceBergen Commitment to Compliance

- Each of ABC's 38 Distribution Centers (DCs) have a Compliance Coordinator responsible for monitoring adherence to state and federal regulations and company policy & procedures.

- **Corporate Support Function:**
  - Regulatory Affairs Supervisor for each region
  - Audit each Distribution Center regularly
  - Annual Compliance Training Conference
  - Periodic Alerts & Updates
  - Quarterly Newsletter
  - Internal Website

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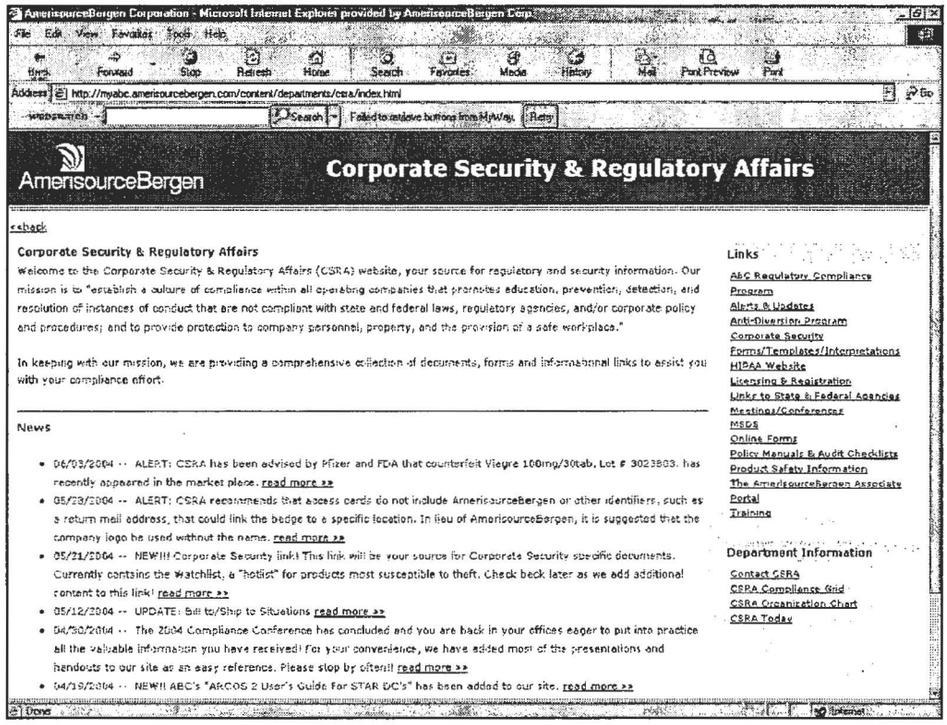

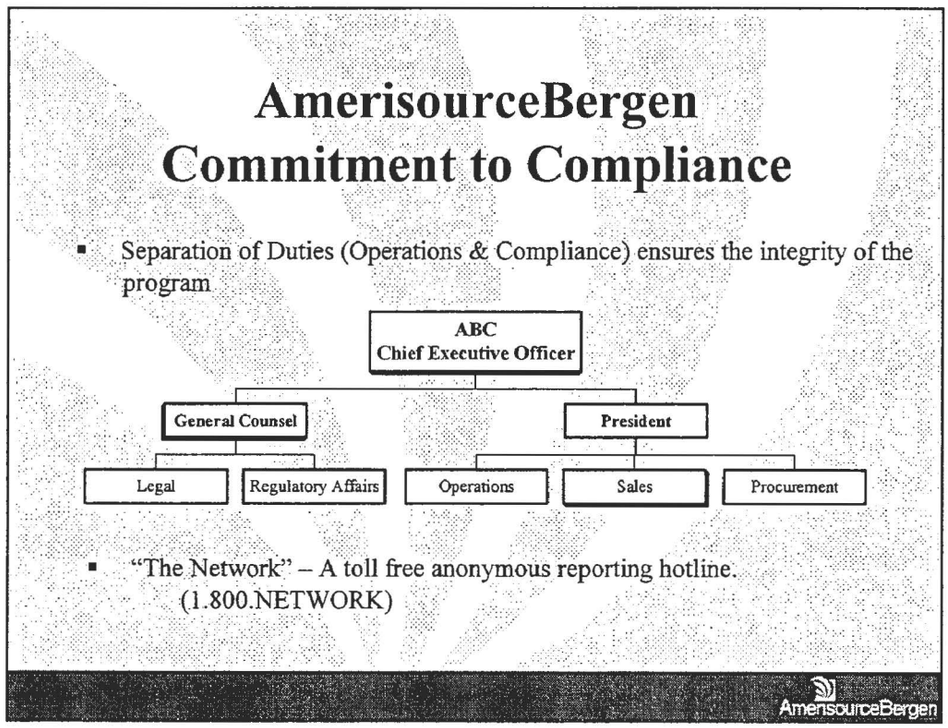

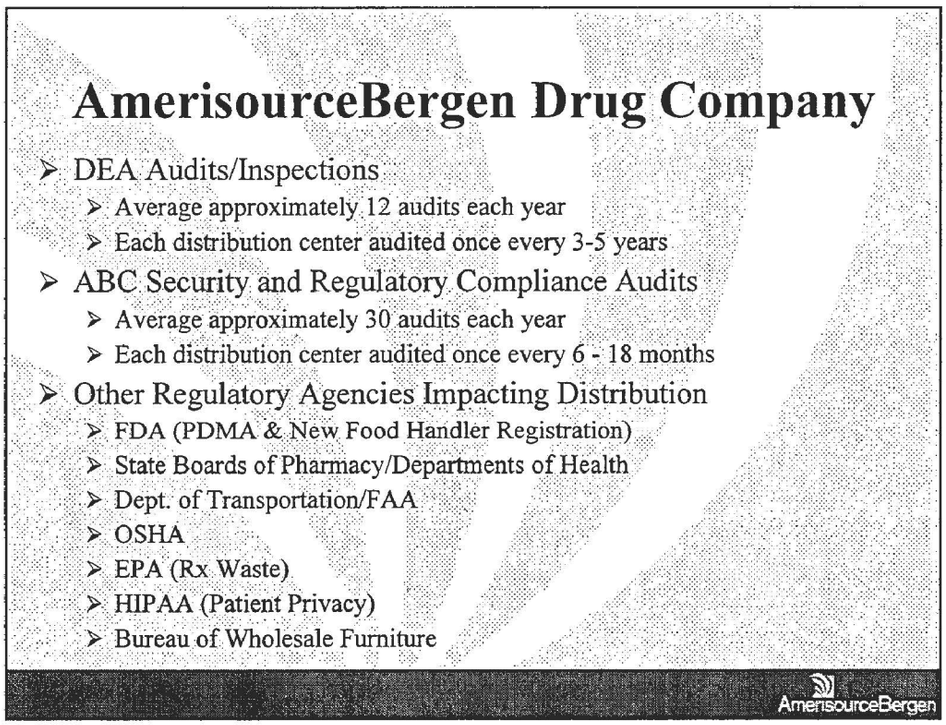

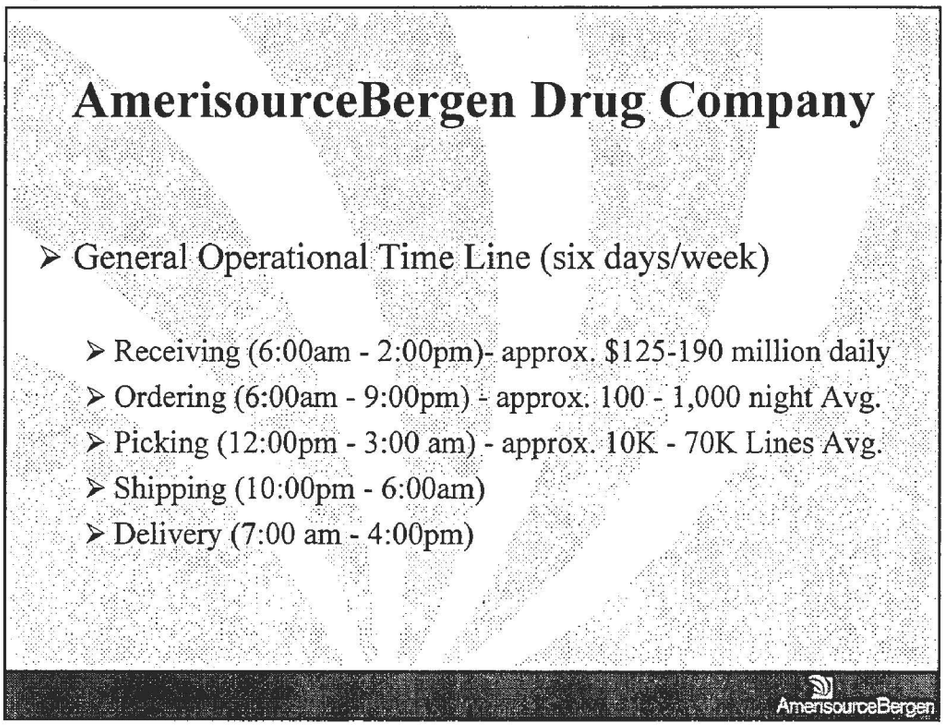

Standardization

- All ABC DCs operate under the same procedures.
- ABC DC security systems & policies were developed to comply with Title 21 of the Code of Federal Regulations:
  - 21 CFR, Part 1300 – Storage & Distribution of Controlled Substances & Listed Chemicals
  - 21CFR, Part 200 – Storage & Distribution of pharmaceutical product (PDMA)

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Code of Federal Regulations

- Interpretation
  - Between different facets of the Industry (Phcy, Dist, Mfg's)
  - Between Industry and DEA
  - Between DEA Field Offices and/or Washington

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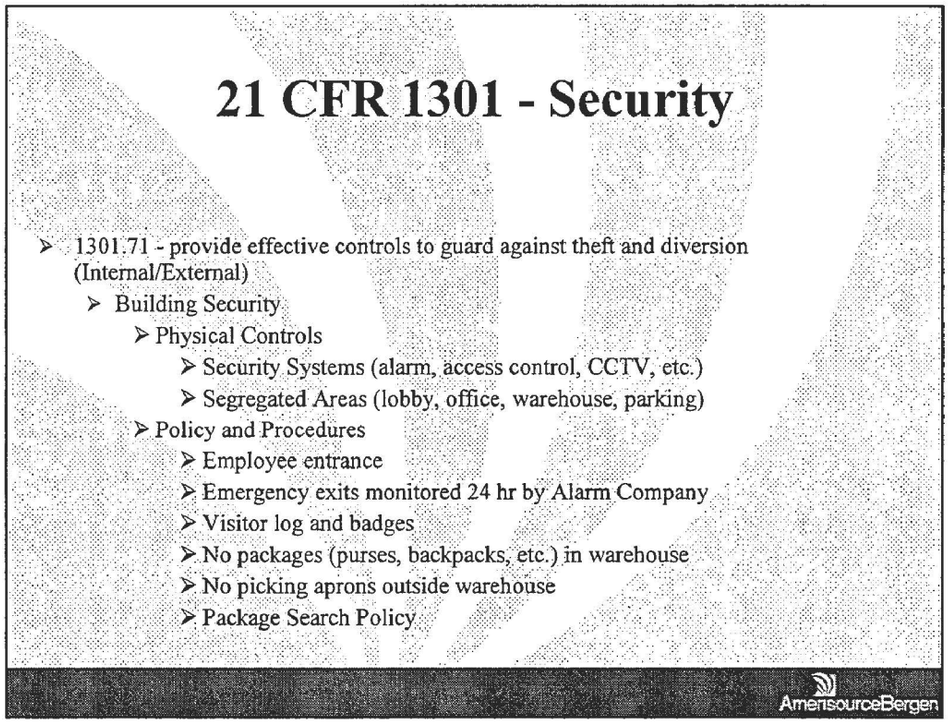

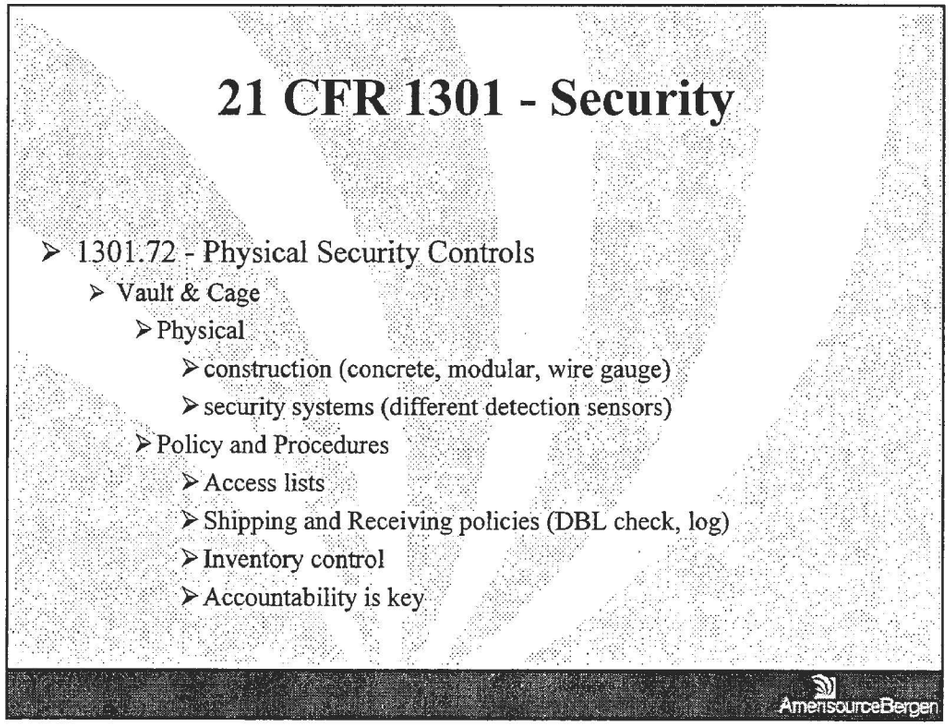

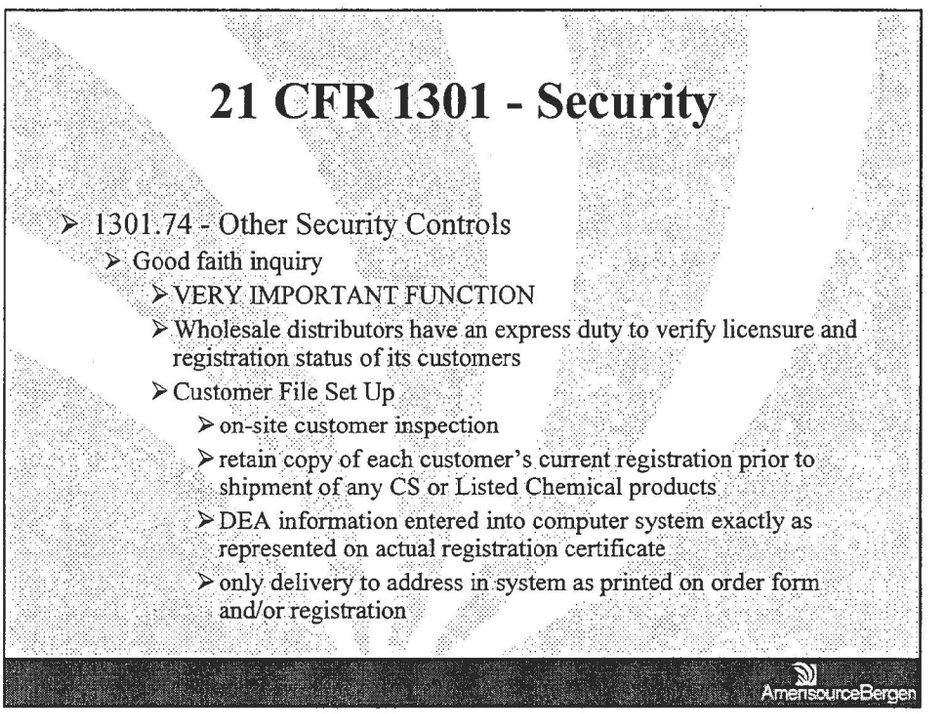

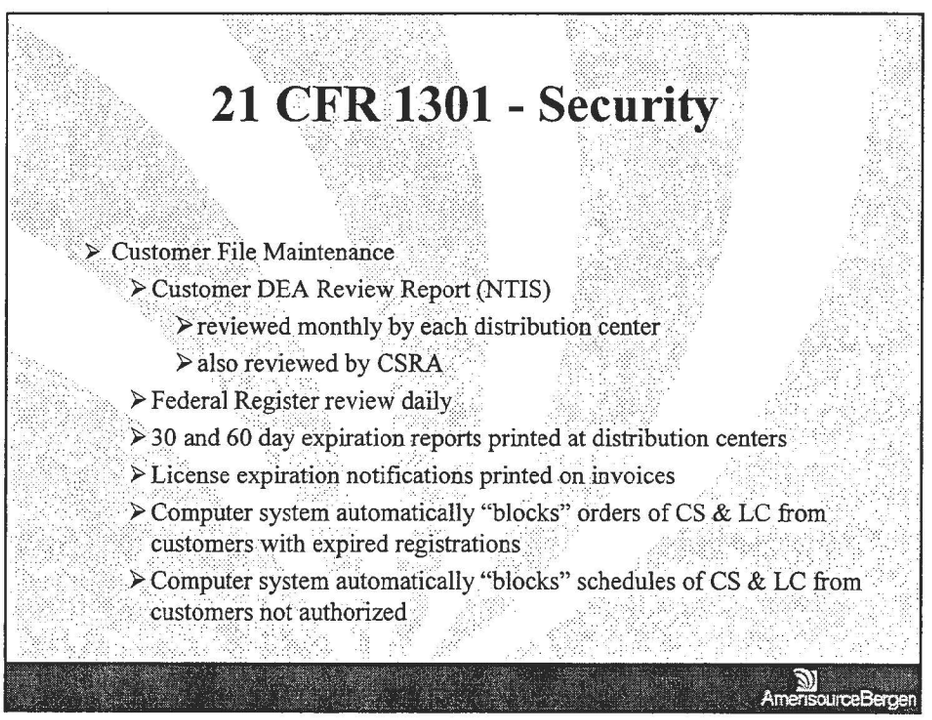

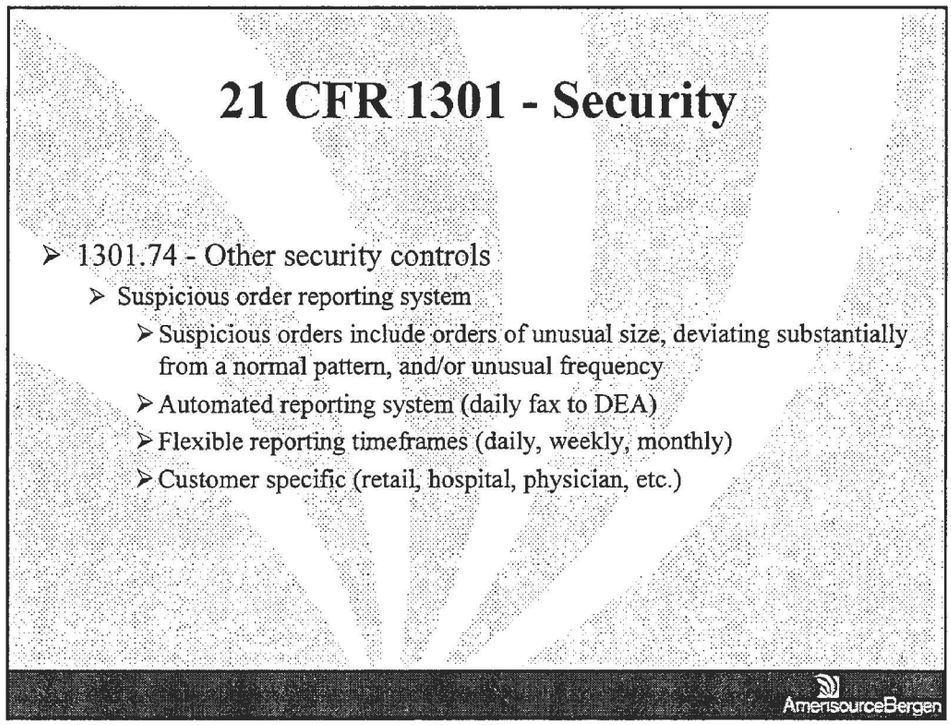

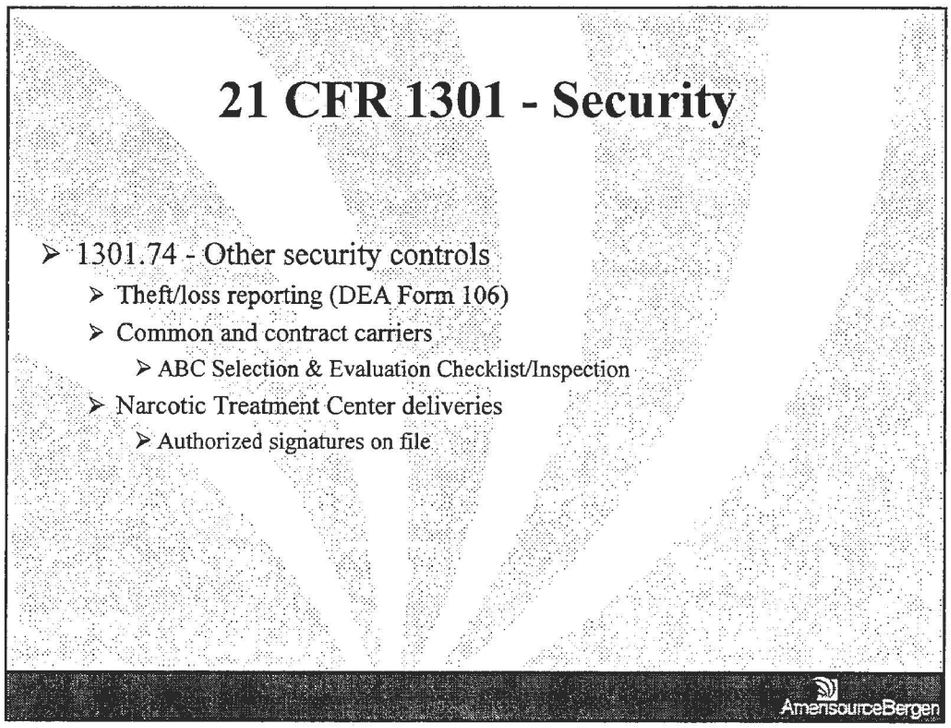

21 CFR 1301 - Security

- 1301.90 & .93 - Employee Screening
  - Prospective employees including temporary associates
  - **All Associates**
    - "Justifacts" - ALL felony & misdemeanor convictions
    - Drug Screens – Initial and Random
  - **"Compliance Critical" Personnel**
    - DEA - Drug related charges, denials, revocations, etc.
    - Annual "Justifacts" criminal record checks

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21 CFR 1304 – Records & Reports

- 1304.04 - Maintenance of records and inventories
  - 2 years (3 years maintained per PDMA guidelines)
  - Central Recordkeeping
    - not including executed DEA Form 222's or inventories
  - Records must be made available within two business days of a written request
  - Schedule II records must be stored separately
  - Schedule III-V records stored separately or in a readily retrievable manner

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21 CFR 1304 – Records & Reports

- 1304.11 - Inventory requirements
  - Complete and accurate record of all controlled substances on hand on the date the inventory is taken
  - ABC Policy and Procedures regarding shipping, receiving and returns and accurate dates
  - Inventories
    - Daily activity counts (detect and correct errors)
    - Monthly inventory of all Controlled Substances on hand in the building (saleable & unsaleable)
    - Year End ARCOS - Schedule II & III narcotics (Dec. 31st COB)
    - Two complete physical inventories of all product annually.
    - Biennial - a complete inventory within two years of Initial or last Biennial inventory (ABC - Dec. 31st every even-numbered year)

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21 CFR 1304 – Records & Reports

- Area of confusion during audits/inspections
- 1304.22 - Records for distributors
  - Product name, description, quantity, name, addresses, DEA registration number, for all transactions (sales, receipt, credit, debit, etc.)
  - ABC Invoice & Credit Memo (See samples)
  - All of this information is available
- 1304.33 Reports to ARCOS
  - Filed monthly
  - All Schedule II and Schedule III narcotic products

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21 CFR 1305 - Order Forms

- 1305.03, .06, .09 & .11 - An order form (DEA Form 222) is required for each distribution of a Schedule II controlled substance
  - Alterations, incomplete forms
  - Each DEA Form 222 is double-checked and matched against work order for accuracy and completeness
- 1305.07 - Power of Attorney
  - Authorizes other associates to sign DEA Form 222's
- 1305.15 - Cancellations
  - Purchaser cancellations
  - Supplier cancellations
- Automation (CSOS) will positively impact the Industry

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21 CFR 1309 – Listed Chemicals

- Registration, Recordkeeping and Reporting
- 1309.21 - (Registration) Every person who distributes List I Chemicals shall obtain annually a registration specific to List I Chemicals…..
  - Exemptions
    - DEA Controlled Substance registration
    - "retail distributor"
- 1309.23 - Separate registration for separate locations
  - System automatically blocks customers without registration

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